Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number UK MIA(IMP) 18693

2: Name of authorisation holder FISHER CLINICAL SERVICES UK LIMITED

FISHER CLINICAL SERVICES UK LIMITED,

LANGHURSTWOOD ROAD, HORSHAM, RH12 4QD, UNITED

KINGDOM

FISHER CLINICAL SERVICES UK LIMITED,

LANGHURSTWOOD ROAD, HORSHAM, RH12 4QD, UNITED

KINGDOM

5: Scope of authorisation and dosage forms ANNEX 1 and/ or ANNEX 2

Part 6 of The Medicines for Human Use (Clinical Trials)

Regulations 2004 [SI 2004/1031]

7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation

member state granting the mandacturing authorisat

4: Legally registered address of authorisation holder

29/05/2025

Confidential

8: Authorisation Date
9: Annexes attached

6: Legal Basis of authorisation

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

3: Address(es) of manufacturing site(s)

Annex 2

Name and address of the site:

FISHER CLINICAL SERVICES UK LIMITED, LANGHURSTWOOD ROAD, HORSHAM, RH12 4QD, UNITED KINGDOM

Human Investigational Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

Part 1 - MANUFACTURING OPERATIONS

[1.1] Sterile Investigational Medicinal Products

[1.1.3] Batch certification

[1.2] Non-sterile investigational medicinal products

[1.2.2] Batch certification

[1.3] Biological investigational medicinal products

[1.3.2] Batch certification

[1.3.2.1] Blood products

Issue Date: 29 May 2025

[1.3.2.2] Immunological products [1.3.2.3] Cell therapy products [1.3.2.4] Gene therapy products [1.3.2.5] Biotechnology products [1.3.2.6] Human or animal extracted products [1.3.2.8] Other biological medicinal products **IMPs** [1.5] Packaging [1.5.1] Primary packaging [1.5.1.1] Capsules, hard shell [1.5.1.2] Capsules, soft shell [1.5.1.8] Other solid dosage forms [1.5.1.13] Tablets [1.5.2] Secondary packaging Part 2 - IMPORTATION OF MEDICINAL PRODUCTS [2.2] Batch certification of imported medicinal products [2.2.1] Sterile Products [2.2.1.1] Aseptically prepared [2.2.1.2] Terminally sterilised [2.2.2] Non-sterile products [2.2.3] Biological medicinal products [2.2.3.1] Blood products [2.2.3.2] Immunological products [2.2.3.3] Cell therapy products [2.2.3.4] Gene therapy products [2.2.3.5] Biotechnology products [2.2.3.6] Human or animal extracted products [2.3] Other Importation Activities [2.3.1] Site of Physical Importation



Importation of QP certified IMPs from a country on the 'approved country for import list'

[2.3.4] Other

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